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* * * * * * * * * * Welcome to STN International * * * * * * * * * *

NEWS 1		Web Page for STN Seminar Schedule - N. America
NEWS 2	JAN 08	CHEMLIST enhanced with New Zealand Inventory of Chemicals
NEWS 3	JAN 16	CA/CAplus Company Name Thesaurus enhanced and reloaded
NEWS 4	JAN 16	IPC version 2007.01 thesaurus available on STN
NEWS 5	JAN 16	WPIDS/WPINDEX/WPIX enhanced with IPC 8 reclassification data
NEWS 6	JAN 22	CA/CAplus updated with revised CAS roles
NEWS 7	JAN 22	CA/CAplus enhanced with patent applications from India
NEWS 8	JAN 29	PHAR reloaded with new search and display fields
NEWS 9	JAN 29	CAS Registry Number crossover limit increased to 300,000 in multiple databases
NEWS 10	FEB 15	PATDPASPC enhanced with Drug Approval numbers
NEWS 11	FEB 15	RUSSIAPAT enhanced with pre-1994 records
NEWS 12	FEB 23	KOREAPAT enhanced with IPC 8 features and functionality
NEWS 13	FEB 26	MEDLINE reloaded with enhancements
NEWS 14	FEB 26	EMBASE enhanced with Clinical Trial Number field
NEWS 15	FEB 26	TOXCENTER enhanced with reloaded MEDLINE
NEWS 16	FEB 26	IFICDB/IFIPAT/IFIUDB reloaded with enhancements
NEWS 17	FEB 26	CAS Registry Number crossover limit increased from 10,000 to 300,000 in multiple databases
NEWS 18	MAR 15	WPIDS/WPIX enhanced with new FRAGHITSTR display format
NEWS 19	MAR 16	CASREACT coverage extended
NEWS 20	MAR 20	MARPAT now updated daily
NEWS 21	MAR 22	LWPI reloaded
NEWS 22	MAR 30	RDISCLOSURE reloaded with enhancements
NEWS 23	APR 02	JICST-EPLUS removed from database clusters and STN
NEWS 24	APR 30	GENBANK reloaded and enhanced with Genome Project ID field
NEWS 25	APR 30	CHEMCATS enhanced with 1.2 million new records
NEWS 26	APR 30	CA/CAplus enhanced with 1870-1889 U.S. patent records
NEWS 27	APR 30	INPADOC replaced by INPADOCDB on STN
NEWS 28	MAY 01	New CAS web site launched
NEWS 29	MAY 08	CA/CAplus Indian patent publication number format defined
NEWS 30	MAY 14	RDISCLOSURE on STN Easy enhanced with new search and display fields
NEWS 31	MAY 21	BIOSIS reloaded and enhanced with archival data
NEWS 32	MAY 21	TOXCENTER enhanced with BIOSIS reload
NEWS 33	MAY 21	CA/CAplus enhanced with additional kind codes for German patents
NEWS 34	MAY 22	CA/CAplus enhanced with IPC reclassification in Japanese patents

NEWS EXPRESS NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.

```
 NEWS HOURS     STN Operating Hours Plus Help Desk Availability
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Enter NEWS followed by the item number or name to see news on that
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FILE 'HOME' ENTERED AT 11:22:38 ON 13 JUN 2007

=> file reg
COST IN U.S. DOLLARS                                    SINCE FILE      TOTAL
                                                           ENTRY    SESSION
FULL ESTIMATED COST                                         0.21       0.21

FILE 'REGISTRY' ENTERED AT 11:22:46 ON 13 JUN 2007
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STRUCTURE FILE UPDATES:   12 JUN 2007  HIGHEST RN 937161-92-7
DICTIONARY FILE UPDATES:  12 JUN 2007  HIGHEST RN 937161-92-7

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TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

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experimental property data in the original document.  For information
on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

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Uploading C:\Program Files\Stnexp\Queries\10527762\Struc 1.str
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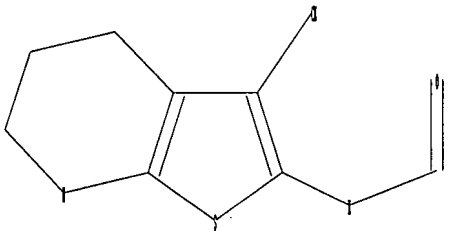

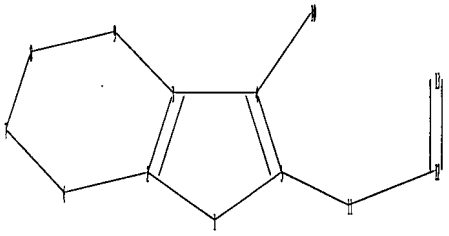

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chain nodes :

10527762.trn
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chain bonds :
4-10  5-11  11-12  12-13
ring bonds :
1-2  1-5  2-3  2-6  3-4  3-9  4-5  6-7  7-8  8-9
exact/norm bonds :
1-2  1-5  2-3  2-6  3-4  3-9  4-5  5-11  6-7  7-8  8-9  11-12  12-13
exact bonds :
4-10


Match level :
1:Atom  2:Atom  3:Atom  4:Atom  5:Atom  6:Atom  7:Atom  8:Atom  9:Atom  10:CLASS
11:CLASS  12:CLASS  13:CLASS



L1      STRUCTURE UPLOADED

=> d
L1 HAS NO ANSWERS
L1                STR
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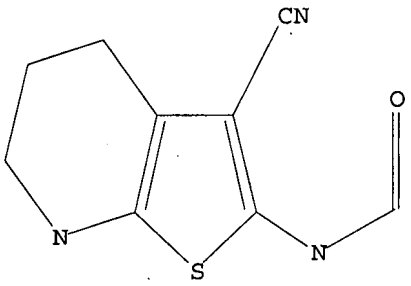


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Structure attributes must be viewed using STN Express query preparation.


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100.0% PROCESSED       0 ITERATIONS                                0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:            0 TO        0
PROJECTED ANSWERS:               0 TO        0

L2             0 SEA SSS SAM L1

=> l1 full
FULL SEARCH INITIATED 11:23:02 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -       54 TO ITERATE

100.0% PROCESSED       54 ITERATIONS                              20 ANSWERS
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SEARCH TIME: 00.00.01

L3 20 SEA SSS FUL L1

=> file medline caplus chemcat

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	172.10	172.31

FILE 'MEDLINE' ENTERED AT 11:23:11 ON 13 JUN 2007

FILE 'CAPLUS' ENTERED AT 11:23:11 ON 13 JUN 2007
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FILE 'CHEMCATS' ENTERED AT 11:23:11 ON 13 JUN 2007
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=> l3
L4 11 L3

=> dup rem l4
DUPLICATE IS NOT AVAILABLE IN 'CHEMCATS'.

ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE
PROCESSING COMPLETED FOR L4
L5 11 DUP REM L4 (0 DUPLICATES REMOVED)

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=> d ibib abs hitstr 1-11
NO VALID FORMATS ENTERED FOR FILE 'CHEMCATS'
In a multifile environment, each file must have at least one valid
format requested.  Refer to file specific help messages or the
STNGUIDE file for information on formats available in individual
files.
REENTER DISPLAY FORMAT FOR ALL FILES (FILEDEFAULT):
REENTER DISPLAY FORMAT FOR ALL FILES (FILEDEFAULT):filedefault
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L5 ANSWER 1 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2007:188037 CAPLUS
DN 146:350594
TI N-(3-Cyano-4,5,6,7-tetrahydro-1-benzothien-2-yl)amides as potent, selective, inhibitors of JNK2 and JNK3
AU Angell, Richard M.; Atkinson, Francis L.; Brown, Murray J.; Chuang, Tsu Tshen; Christopher, John A.; Cichy-Knight, Maria; Dunn, Allison K.; Hightower, Kendra E.; Malkakorpi, Susanna; Musgrave, James R.; Neu, Margarete; Rowland, Paul; Shea, Robyn L.; Smith, Jeffery L.; Somers, Donald O.; Thomas, Sonia A.; Thompson, Gladstone; Wang, Ruolan
CS GlaxoSmithKline R&D, Medicines Research Centre, Stevenage, Hertfordshire, SG1 2NY, UK
SO Bioorganic & Medicinal Chemistry Letters (2007), 17(5), 1296-1301
CODEN: BMCLE8; ISSN: 0960-894X
PB Elsevier Ltd.
DT Journal
LA English
RE.CNT 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2006:79076 CAPLUS
DN 144:170973
TI Preparation of (fused) thienopyridines for treatment of hepatitis C infection.
IN Karp, Gary Mitchell; Chen, Guangming
PA USA
SO U.S. Pat. Appl. Publ., 186 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2006019976	A1	20060126	US 2005-180779	20050714
	AU 2005275182	A1	20060223	AU 2005-275182	20050714
	CA 2578636	A1	20060223	CA 2005-2578636	20050714
	WO 2006019832	A1	20060223	WO 2005-US24882	20050714
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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PRAI	US 2004-589876P	P	20040722		
	WO 2005-US24882	W	20050714		

OS MARPAT 144:170973

L5 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2005:423698 CAPLUS
DN 142:458555
TI Preparation of 2-aminothiophene derivatives as fungicides
IN Selles, Patrice; Wailes, Jeffrey Steven; Whittingham, William Guy; Clarke, Eric Daniel
PA Syngenta Participations A.-G., Switz.; Syngenta Limited
SO PCT Int. Appl., 155 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

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PI	WO 2005044008	A2	20050519	WO 2004-GB4429	20041019
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI	GB 2003-24653	A	20031022		

OS MARPAT 142:458555

L5 ANSWER 4 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 2004:252284 CAPLUS
DN 140:287368
TI Preparation of fused thiophenes as glucagon receptor blockers for treatment of type 2 diabetes.
IN Duffy, Joseph; Campbell, Elizabeth Louise; Liang, Rui; Konteatis, Zenon
PA Merck & Co., Inc., USA
SO PCT Int. Appl., 47 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004024065	A2	20040325	WO 2003-US28033	20030908
	WO 2004024065	A3	20040513		
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	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2498106	A1	20040325	CA 2003-2498106	20030908
	AU 2003270390	A1	20040430	AU 2003-270390	20030908
	EP 1549655	A2	20050706	EP 2003-752080	20030908
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	JP 2006503034	T	20060126	JP 2004-536137	20030908
	US 2005239865	A1	20051027	US 2005-527762	20050311
PRAI	US 2002-410145P	P	20020912		
	WO 2003-US28033	W	20030908		

OS MARPAT 140:287368

L5 ANSWER 5 OF 11 CAPLUS COPYRIGHT 2007 ACS on STN
AN 1972:526675 CAPLUS
DN 77:126675
TI Antiviral 5,6,7,8-tetrahydro-5,8-ethanopyridino[2,3-b]thieno[5,4-d]pyrimidines
IN Wellings, Ian
SO U.S., 7 pp.
CODEN: USXXAM
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 3681351	A	19720801	US 1970-28959	19700415
PRAI US 1970-28959	A	19700415		

L5 ANSWER 6 OF 11 CHEMCATS COPYRIGHT 2007 ACS on STN
Accession No. (AN): 2007:2783940 CHEMCATS
Catalog Name (CO): Chemical Block Stock Library
Publication Date (PD): 24 May 2007
Order Number (ON): A4016/0171272
Chemical Name (CN): Urea, N-(3-cyano-5,6-dihydro-4H-4,7-ethanothieno[2,3-b]pyridin-2-yl)-N'-(3,4-dichlorophenyl)-
CAS Registry No. (RN): 874590-25-7
Supplementary Term (ST): CHEMICAL LIBRARY
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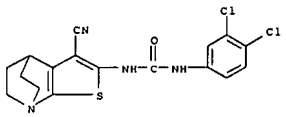

L5 ANSWER 7 OF 11 CHEMCATS COPYRIGHT 2007 ACS on STN
Accession No. (AN): 2007:1841359 CHEMCATS
Catalog Name (CO): Ambinter Stock Screening Collection
Publication Date (PD): 15 Feb 2007
Order Number (ON): A4016/0171272
Chemical Name (CN): Chemical name not yet assigned
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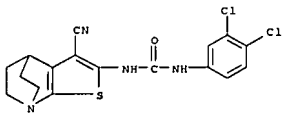

L5 ANSWER 8 OF 11 CHEMCATS COPYRIGHT 2007 ACS on STN
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Catalog Name (CO): Scientific Exchange Product List
Publication Date (PD): 18 May 2007
Order Number (ON): M-919798
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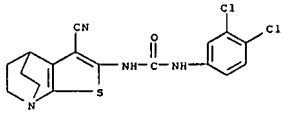

L5 ANSWER 9 OF 11 CHEMCATS COPYRIGHT 2007 ACS on STN
Accession No. (AN): 2006:3038777 CHEMCATS
Catalog Name (CO): MicroChemistry Screening Collection
Publication Date (PD): 25 Apr 2007
Order Number (ON): 239400
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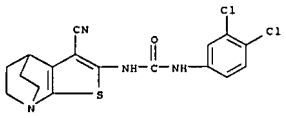

L5 ANSWER 10 OF 11 CHEMCATS COPYRIGHT 2007 ACS on STN
Accession No. (AN): 2006:1652634 CHEMCATS
Catalog Name (CO): Aurora Screening Library
Publication Date (PD): 1 Jan 2007
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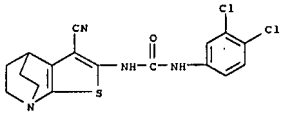

L5 ANSWER 11 OF 11 CHEMCATS COPYRIGHT 2007 ACS on STN
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Catalog Name (CO): AKos Screening Library
Publication Date (PD): 7 Feb 2006
Order Number (ON): AKLM5-PFR-150818
Chemical Name (CN): Chemical name not yet assigned
CAS Registry No. (RN): 874590-25-7
Supplementary Term (ST): CHEMICAL LIBRARY
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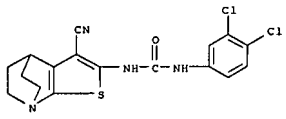

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COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                         ENTRY    SESSION
FULL ESTIMATED COST                                      19.97     192.28

FILE 'MEDLINE' ENTERED AT 11:24:18 ON 13 JUN 2007

FILE 'CAPLUS' ENTERED AT 11:24:18 ON 13 JUN 2007
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=> l3
L6            5 L3

=> d ibib abs hitstr 1-5
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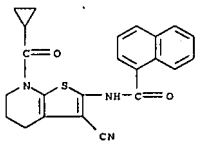
L6 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2007:188037 CAPLUS
DOCUMENT NUMBER: 146:350594
TITLE:
N-(3-Cyano-4,5,6,7-tetrahydro-1-benzothien-2-yl)amides as potent, selective, inhibitors of JNK2 and JNK3
AUTHOR(S): Angell, Richard M.; Atkinson, Francis L.; Brown, Murray J.; Chuang, Tsu Tshen; Christopher, John A.; Cichy-Knight, Maria; Dunn, Allison K.; Hightower, Kendra E.; Malkakorpi, Susanna; Musgrave, James R.; Neu, Margarete; Rowland, Paul; Shea, Robyn L.; Smith, Jeffery L.; Somers, Donald O.; Thomas, Sonia A.; Thompson, Gladstone; Wang, Ruolan
CORPORATE SOURCE: GlaxoSmithKline R&D, Medicines Research Centre, Stevenage, Hertfordshire, SG1 2NY, UK
SOURCE: Bioorganic & Medicinal Chemistry Letters (2007), 17(5), 1296-1301
CODEN: BMCLE8; ISSN: 0960-894X
PUBLISHER: Elsevier Ltd.
DOCUMENT TYPE: Journal
LANGUAGE: English

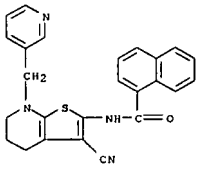
AB The identification and exploration of a novel, potent and selective series of N-(3-cyano-4,5,6,7-tetrahydro-1-benzothien-2-yl)amide inhibitors of JNK2 and JNK3 kinases is described. Compds. 5a and 11a were identified as potent inhibitors of JNK3 (pIC50 6.7 and 6.6, resp.), with essentially equal potency against JNK2 (pIC50 6.5). Selectivity within the mitogen-activated protein kinase (MAPK) family, against JNK1, p38α and ERK2, was observed for the series. X-ray crystallog. of 5e and 8a in JNK3 revealed a unique binding mode, with the 3-cyano substituent forming an H-bond acceptor interaction with the hinge region of the ATP-binding site.

IT 929700-76-5P 929700-77-6P 929700-78-7P
929700-79-8P 929700-80-1P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(cyanotetrahydrobenzothienylamides as inhibitors of JNK2 and JNK3)

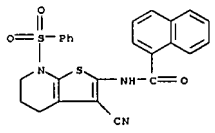
RN 929700-76-5 CAPLUS
CN 1-Naphthalenecarboxamide, N-[3-cyano-7-(cyclopropylcarbonyl)-4,5,6,7-tetrahydrothieno[2,3-b]pyridin-2-yl]- (CA INDEX NAME)



L6 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

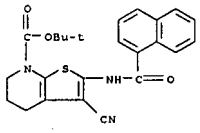


RN 929700-80-1 CAPLUS
CN 1-Naphthalenecarboxamide, N-[3-cyano-4,5,6,7-tetrahydro-7-(phenylsulfonyl)thieno[2,3-b]pyridin-2-yl]- (CA INDEX NAME)



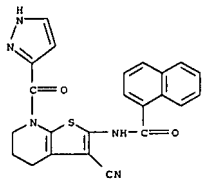
IT 929700-66-3P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(cyanotetrahydrobenzothienylamides as inhibitors of JNK2 and JNK3)

RN 929700-66-3 CAPLUS
CN Thieno[2,3-b]pyridine-7(4H)-carboxylic acid, 3-cyano-5,6-dihydro-2-[(1-naphthalenylcarbonyl)amino]-, 1,1-dimethylethyl ester (CA INDEX NAME)

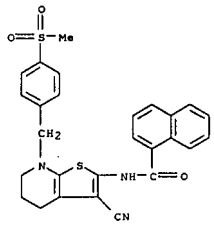


REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)
RN 929700-77-6 CAPLUS
CN 1-Naphthalenecarboxamide, N-(3-cyano-4,5,6,7-tetrahydro-7-(1H-pyrazol-3-ylcarbonyl)thieno[2,3-b]pyridin-2-yl]- (CA INDEX NAME)



RN 929700-78-7 CAPLUS
CN 1-Naphthalenecarboxamide, N-[3-cyano-4,5,6,7-tetrahydro-7-[[4-(methylsulfonyl)phenyl]methyl]thieno[2,3-b]pyridin-2-yl]- (CA INDEX NAME)



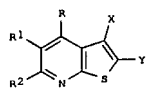
RN 929700-79-8 CAPLUS
CN 1-Naphthalenecarboxamide, N-[3-cyano-4,5,6,7-tetrahydro-7-(3-pyridinylmethyl)thieno[2,3-b]pyridin-2-yl]- (CA INDEX NAME)

L6 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2006:79076 CAPLUS
DOCUMENT NUMBER: 144:170973
TITLE: Preparation of (fused) thienopyridines for treatment of hepatitis C infection.
INVENTOR(S): Karp, Gary Mitchell; Chen, Guangming
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 186 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2006019976	A1	20060126	US 2005-180779	20050714
AU 2005275182	A1	20060223	AU 2005-275182	20050714
CA 2578636	A1	20060223	CA 2005-2578636	20050714
WO 2006019832	A1	20060223	WO 2005-US24882	20050714
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
EP 1781289	A1	20070509	EP 2005-773284	20050714
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				

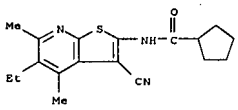
PRIORITY APPLN. INFO.: US 2004-589876P P 20040722
WO 2005-US24882 W 20050714

OTHER SOURCE(S): MARPAT 144:170973
GI

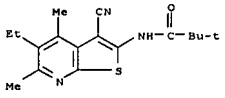


AB Title compds. [I; X = H, cyano, amino, heteroaryl, alkoxy, cyano, halo, etc.; Y = halo, amino, alkylsulfonyl, cyano, (substituted) aryl, amino, heterocyclyl, heteroaryl, aryl, etc.; R = H, alkyl, haloalkyl, hydroxyalkyl, aryl, haloaryl; R1 = H, aryl, alkyl, alkoxy, aminoalkoxy, heterocyclylalkoxy, amino, etc.; R2 = alkyl, heterocyclyl, amino; adjacent

L6 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)
pairs of variables may form rings], were prepd. Thus, 2-cyanothioacetamide, 3-ethylpentane-2,4-dione, and Et3N were heated in EtOH at 60° for 1 h to give 89% 5-ethyl-2-mercapto-4,6-dimethylnicotinonitrile. This was stirred with tert-Bu bromoacetate and K2CO3 in DMF at room temp. to 80° to give 96% tert-Bu 3-amino-5-ethyl-4,6-dimethylnicotinonitrile. Several I showed IC50's of <0.5 μM in an HCV replicon system.
IT 874633-06-4P 874633-07-5P 874633-08-6P
874633-13-3P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(claimed compound; preparation of (fused) thienopyridines for treatment of
hepatitis C infection)
RN 874633-06-4 CAPLUS
CN Cyclopentanecarboxamide, N-(3-cyano-5-ethyl-4,6-dimethylthieno[2,3-b]pyridin-2-yl)- (9CI) (CA INDEX NAME)

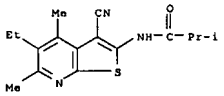


RN 874633-07-5 CAPLUS
CN Propanamide, N-(3-cyano-5-ethyl-4,6-dimethylthieno[2,3-b]pyridin-2-yl)-2,2-dimethyl- (9CI) (CA INDEX NAME)

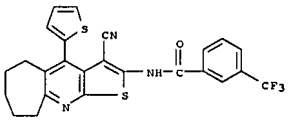


RN 874633-08-6 CAPLUS
CN Propanamide, N-(3-cyano-5-ethyl-4,6-dimethylthieno[2,3-b]pyridin-2-yl)-2-methyl- (9CI) (CA INDEX NAME)

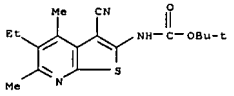
L6 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)



RN 874633-13-3 CAPLUS
CN Benzamide, N-[3-cyano-6,7,8,9-tetrahydro-4-(2-thienyl)-5H-cyclohepta[b]thieno[3,2-e]pyridin-2-yl]-3-(trifluoromethyl)- (9CI) (CA INDEX NAME)



IT 874633-38-2P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of (fused) thienopyridines for treatment of hepatitis C infection)
RN 874633-38-2 CAPLUS
CN Carbamic acid, (3-cyano-5-ethyl-4,6-dimethylthieno[2,3-b]pyridin-2-yl)-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



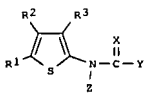
L6 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2005:423698 CAPLUS
DOCUMENT NUMBER: 142:458555
TITLE: Preparation of 2-aminothiophene derivatives as fungicides
INVENTOR(S): Selles, Patrice; Wailes, Jeffrey Steven; Whittingham, William Guy; Clarke, Eric Daniel
PATENT ASSIGNEE(S): Syngenta Participations A.-G., Switz.; Syngenta Limited
SOURCE: PCT Int. Appl., 155 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005044008	A2	20050519	WO 2004-GB4429	20041019

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

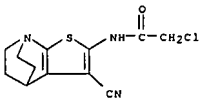
PRIORITY APPLN. INFO.: GB 2003-24653 A 20031022

OTHER SOURCE(S): MARPAT 142:458555
GI



AB The 2-aminothiophene derivs. I [R1, R2 = H, halo, (cyclo)alkyl, hydroxyalkyl, etc.; R1R2= alkylene; R3 = H, halo, NO2, CN, (halo)alkyl, alkenyl, alkynyl, etc.; X = O, S, NH2,etc.; Y = H, (halo)alkyl, hydroxyalkyl, etc.; Z = H, (alkoxy)alkyl, alkylcarbonyl, etc.] are prepared
as fungicides. The invention further relates to fungicidal compns. containing
these compds., processes for preparing these compds. and to some of the compds. themselves.
IT 851143-96-4P
RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation as fungicide)
RN 851143-96-4 CAPLUS
CN Acetamide, 2-chloro-N-(3-cyano-5,6-dihydro-4H-4,7-ethanothieno[2,3-

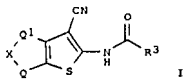
L6 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)
b]pyridin-2-yl)- (9CI) (CA INDEX NAME)



L6 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2004:252284 CAPLUS
DOCUMENT NUMBER: 140:287368
TITLE: Preparation of fused thiophenes as glucagon receptor blockers for treatment of type 2 diabetes.
INVENTOR(S): Duffy, Joseph; Campbell, Elizabeth Louise; Liang, Rui; Konteatis, Zenon
PATENT ASSIGNEE(S): Merck & Co., Inc., USA
SOURCE: PCT Int. Appl., 47 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004024065	A2	20040325	WO 2003-US28033	20030908
WO 2004024065	A3	20040513		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2498106	A1	20040325	CA 2003-2498106	20030908
AU 2003270390	A1	20040430	AU 2003-270390	20030908
EP 1549655	A2	20050706	EP 2003-752080	20030908
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2006503034	T	20060126	JP 2004-536137	20030908
US 2005239865	A1	20051027	US 2005-527762	20050311
PRIORITY APPLN. INFO.:			US 2002-410145P	P 20020912
			WO 2003-US28033	W 20030908

OTHER SOURCE(S): MARPAT 140:287368
GI

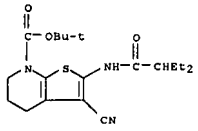


I

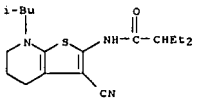
AB Title compds. [I; Q = (CR5R6)m; Q1 = (CH2)n; X = NR4, CR5R6; R1 = H, (substituted) alkyl, cycloalkyl, aryl; R2 = R1, CO2R7, CONR7R8; m, n = 0-3; R3 = (substituted) alkyl, cycloalkyl, aryl; R4 = (substituted) alkyl, cycloalkyl, aryl, arylcarbonyl, heteroaryl, heteroarylcarbonyl, etc.; 1 of R5, R6 = NR11R12, NR11COR12, NR11CO2R12, NR11SO2R12, the other = R1, OR11, heteroaryl, etc.; R7, R10, R11 = R1, (substituted) heteroaryl, etc.; R8, R12 = (substituted) alkyl, cycloalkyl, aryl, heteroaryl, etc.; R11R12 = atoms to form a 5-8 membered (substituted) ring; with provisos], were prepd. for treatment of diabetes and related conditions (no data). Thus, tert-Bu 3-oxopiperidine-1-carboxylate, malononitrile, morpholine, and S were stirred 16 h in EtOH to give tert-Bu 2-amino-3-cyano-5,6-dihydrothieno[2,3-b]pyridine-7(4H)-carboxylate. This was stirred 16 h with diisopropylethylamine and 2-ethylbutanoyl chloride in CH2Cl2 to give tert-Bu 2-[(2-ethylbutanoyl)amino]-3-cyano-5,6-dihydrothieno[2,3-b]pyridine-7(4H)-carboxylate.

IT 675572-21-1P 675572-22-2P 675572-23-3P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(claimed compound; preparation of fused thiophenes as glucagon receptor blockers for treatment of type 2 diabetes)

RN 675572-21-1 CAPLUS
CN Thieno[2,3-b]pyridine-7(4H)-carboxylic acid, 3-cyano-2-[(2-ethyl-1-oxobutyl)amino]-5,6-dihydro-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

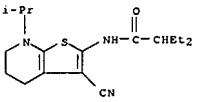


RN 675572-22-2 CAPLUS
CN Butanamide, N-[3-cyano-4,5,6,7-tetrahydro-7-(2-methylpropyl)thieno[2,3-b]pyridin-2-yl]-2-ethyl- (9CI) (CA INDEX NAME)



RN 675572-23-3 CAPLUS
CN Butanamide, N-[3-cyano-4,5,6,7-tetrahydro-7-(1-methylethyl)thieno[2,3-b]pyridin-2-yl]-2-ethyl- (9CI) (CA INDEX NAME)

L6 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

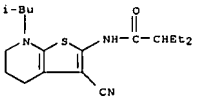


IT 675572-74-4P 675572-75-5P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of fused thiophenes as glucagon receptor blockers for treatment of type 2 diabetes)

RN 675572-74-4 CAPLUS
CN Butanamide, N-[3-cyano-4,5,6,7-tetrahydro-7-(2-methylpropyl)thieno[2,3-b]pyridin-2-yl]-2-ethyl-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

CM 1

CRN 675572-22-2
CMF C18 H27 N3 O S



CM 2

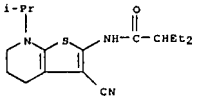
CRN 76-05-1
CMF C2 H F3 O2

RN 675572-75-5 CAPLUS
CN Butanamide, N-[3-cyano-4,5,6,7-tetrahydro-7-(1-methylethyl)thieno[2,3-b]pyridin-2-yl]-2-ethyl-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

CM 1

CRN 675572-23-3

L6 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)
CMF C17 H25 N3 O S

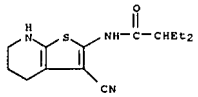


CM 2

CRN 76-05-1
CMF C2 H F3 O2

IT 675572-67-5P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of fused thiophenes as glucagon receptor blockers for treatment of type 2 diabetes)

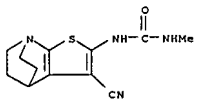
RN 675572-67-5 CAPLUS
CN Butanamide, N-(3-cyano-4,5,6,7-tetrahydrothieno[2,3-b]pyridin-2-yl)-2-ethyl- (9CI) (CA INDEX NAME)



L6 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 1972:526675 CAPLUS
DOCUMENT NUMBER: 77:126675
TITLE: Antiviral 5,6,7,8-tetrahydro-5,8-ethanopyridino[2,3-b]thieno[5,4-d]pyrimidines
INVENTOR(S): Wellings, Ian
SOURCE: U.S., 7 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 3681351	A	19720801	US 1970-28959	19700415
PRIORITY APPLN. INFO.:			US 1970-28959 A	19700415

GI For diagram(s), see printed CA Issue.
AB The title compds. (I, R = H2N, HO, Me2CHNH, HS, H, Cl, MeNH, R1 = H, Me, H2N; and II, R2 = H, Me, EtNH, HO; R3 = H, Me) and their acid salts were prepared by treating III (R4 = cyano, R5 = R6 = H) with (EtO)3CH to give III [R4 = cyano, (R5R6) = :CHOEt (IV) which was aminated by NH3 or secondary amines to give I or II, resp. Thus, 0.1 mole III (R4 = cyano, R5 = R6 = H) was refluxed with 200 ml (EtO)3CH to give IV, which was stirred in NH3-EtOH with continued NH3 sparging to give I (R = H2N, R1 = H) which was converted to the dihydrochloride by HCl-EtOH.
IT 36909-16-7P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 36909-16-7 CAPLUS
CN Urea, N-(3-cyano-5,6-dihydro-4H-4,7-ethanothieno[2,3-b]pyridin-2-yl)-N'-methyl- (9CI) (CA INDEX NAME)



=> log y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	28.55	220.83

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.90	-3.90

STN INTERNATIONAL LOGOFF AT 11:26:33 ON 13 JUN 2007